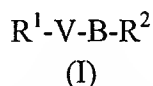


AMENDMENTS TO THE CLAIMS

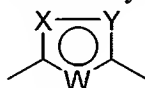
Listing of Claims:

This listing of claims will replace all prior versions and listings of the claims in the application.

1. (Currently Amended) A compound of formula (I), or a pharmaceutically acceptable salt thereof:



wherein V represents a 5-membered heteroaryl ring of the formula:



wherein W is N and one of X and Y is N and the other is O;

B is -CH=CH- or (CH₂)_n, where one of the CH₂ groups may be replaced by O, NR⁵, S(O)_m, C(O) or C(O)NR¹²;

n is 2 or 3;

m is 0, 1 or 2;

R¹ is 4-pyridyl optionally substituted by 1 or 2 halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₃₋₇ cycloalkyl, aryl, OR⁶, CN, NO₂, S(O)_mR⁶, CON(R⁶)₂, N(R⁶)₂, NR¹⁰COR⁶, NR¹⁰SO₂R⁶, SO₂N(R⁶)₂, 4- to 7-membered heterocyclyl or 5- or 6-membered heteroaryl groups;

R² is 4- to 7-membered cycloalkyl substituted by R³, C(O)OR³, C(O)R³ or S(O)₂R³, or R² is 4- to 7-membered heterocyclyl, wherein the heterocycle contains one nitrogen atom which is substituted by~~containing one or two nitrogen atoms which is unsubstituted or substituted by~~ C(O)OR⁴, C(O)R³, S(O)₂R³, C(O)NHR⁴, P(O)(OR¹¹)₂ or a 5- or 6-membered nitrogen containing heteroaryl group;

R³ is C₃₋₈ alkyl, C₃₋₈ alkenyl or C₃₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be optionally substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁴ is C₂₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be substituted with one or more

substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁵ is hydrogen, C(O)R⁷, S(O)₂R⁸, C₃₋₇ cycloalkyl or C₁₋₄ alkyl optionally substituted by OR⁶, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₂ alkyl, C₁₋₂ fluoroalkyl, OR⁶, CN, N(R⁶)₂ and NO₂;

R⁶ are independently hydrogen C₁₋₄ alkyl, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁹, CN, SO₂CH₃, N(R¹⁰)₂ and NO₂; or a group N(R¹⁰)₂ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR¹⁰;

R⁷ is hydrogen, C₁₋₄ alkyl, OR⁶, N(R⁶)₂, aryl or heteroaryl;

R⁸ is C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, aryl or heteroaryl;

R⁹ is hydrogen, C₁₋₂ alkyl or C₁₋₂ fluoroalkyl;

R¹⁰ is hydrogen or C₁₋₄ alkyl;

R¹¹ is phenyl; and

R¹² is hydrogen, C₁₋₄ alkyl or C₃₋₇ cycloalkyl;

provided that the compound is not:

- a) 4-(5-piperidin-4-yl-[1,2,4]oxadiazol-3-yl)pyridine;
- b) 4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid ^tbutyl ester; or
- c) 4-[5-(4-butylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine.

2-8. (Canceled).

9. (Previously Presented) A compound according to claim 1, or a pharmaceutically acceptable salt thereof; wherein R¹ is 4-pyridyl optionally substituted by halo, C₁₋₄ alkyl, C₁₋₄ alkoxy or CN.

10. (Previously Presented) A compound according to claim 1, or a pharmaceutically acceptable salt thereof, wherein R² is a 4- to 7-membered cycloalkyl substituted by R³, or 4- to 7-membered heterocyclyl containing one nitrogen atom which is substituted by C(O)OR⁴.

11. (Previously Presented; Withdrawn) A compound according to claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is C₃₋₈ alkyl which may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl.

12. (Previously Presented) A compound according to claim 1, or a pharmaceutically acceptable salt thereof, wherein R⁴ is C₂₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, 5- to 6-membered heteroaryl containing one or two

nitrogen atoms, C₁₋₄ alkyl, C₃₋₇ cycloalkyl or C₁₋₄ alkylaryl, any of which may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶ and CO₂C₁₋₄ alkyl.

13. **(Original)** A compound according to claim 12, or a pharmaceutically acceptable salt thereof, wherein R⁴ is C₃₋₆ alkyl optionally substituted with up to 5 fluoro or chloro atoms, and which may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl.

14. **(Previously Presented; Withdrawn)** A compound according to claim 1, or a pharmaceutically acceptable salt thereof, wherein R⁵ is C₁₋₄ alkyl.

15. **(Currently Amended)** A compound ~~selected from as defined in any one of Examples 1, 3 to 5, 10 to 13, 16 to 39, 41, 42, 52 to 132, 134, 135, or 147 to 149,~~

4-(3-Pyridin-4-yl)-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid *tert*-butyl ester;

3-(3-Pyridin-4-yl)-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid *tert*-butyl ester;

4-[5-(4-Pentylcyclohexylmethyl)-[1,2,4]oxadiazol-3-yl]pyridine;

trans-2-Chloro-4-[5-(4-pentylcyclohexane)-[1,2,4]oxadiazol-3-yl]pyridine;

4-[5-(4-*n*-Propylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine;

trans-4-[5-(4-Pentylcyclohexane)-[1,2,4]oxadiazol-3-yl]pyridine ;

4-[2-(3-Pyridin-4-yl)-[1,2,4]oxadiazol-5-yl)-ethyl]piperidine-1-carboxylic acid *tert*-butyl ester;

4-(3-Pyridin-4-yl)-[1,2,4]oxadiazol-5-ylmethyl)piperidine-1-carboxylic acid *tert*-butyl ester;

trans-4-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine-2-carboxylic acid methylamide;

trans-4-[5-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine-2-carboxylic acid amide;

trans-4-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-2-Chloro-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-3-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-2-Methyl-3-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-2-Chloro-6-methyl-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-4-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine-2-carbonitrile ;

trans-2-Chloro-3-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-2-Chloro-6-methyl-3-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-2-Methyl-5-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;

trans-3-Methyl-5-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-2,6-Dichloro-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-2-Chloro-6-methoxy-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-5-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]-2-[1,2,4]triazol-1-ylpyridine;
2-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyrazine;
4-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyrimidine;
trans-5-[3-(4-Pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine-2-carbonitrile;
trans-5-Chloro-2-methylsulfanyl-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyrimidine;
trans-2-Fluoro-5-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-2-Fluoro-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-2-Imidazol-1-yl-5-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-2-Methyl-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
trans-3-Methyl-4-[3-(4-pentylcyclohexyl)-[1,2,4]oxadiazol-5-yl]pyridine;
4-(5-Pyridin-4-yl-[1,2,4]oxadiazol-3-ylmethoxy)piperidine-1-carboxylic acid tert-butyl ester;
4-[5-(2-Cyanopyridin-4-yl)-[1,2,4]oxadiazol-3-ylmethoxy]piperidine-1-carboxylic acid tert-butyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid isobutyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid 2-methoxyethyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid ethyl ester;
3,3-Dimethyl-1-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidin-1-yl]butan-1-one;
2-Cyclopentyl-1-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidin-1-yl]ethanone;
4-{5-[1-(Butane-1-sulfonyl)piperidin-4-yl]-[1,2,4]oxadiazol-3-yl}pyridine;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid propylamide;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid tert-butylamide;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid cyclopentyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid benzyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid isobutyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid ethyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid cycloheptyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid methyl ester;

4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2-methoxy-ethyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid isopropyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 4-methoxy-phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2,2,2-trichloroethyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 4-chloro-phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2-ethyl-hexyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid propyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid hexyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid (1R,2S,5R)-2-isopropyl-5-methylcyclohexyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid (1S,2R,5S)-2-isopropyl-5-methylcyclohexyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2,2-dimethylpropyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid naphthalen-1-yl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2-methoxy-phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 3-trifluoromethylphenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid prop-2-ynyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid but-2-ynyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid pentyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid p-tolyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 2-chloro-phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid naphthalen-2-yl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid butyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 4-methoxycarbonyl-phenyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid 4-fluoro-phenyl ester;
3-Methyl-1-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]-butan-1-one;
Phenyl-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]methanone;
1-[4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]butan-1-one;
2,2-Dimethyl-1-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]propan-1-one;

Cyclopentyl-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]methanone;
[4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]-*p*-tolylmethanone;
3,3-Dimethyl-1-[4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]butan-1-one;
4-{5-[1-(Butane-1-sulfonyl) piperidin-4-yloxymethyl]-[1,2,4]oxadiazol-3-yl}pyridine;
4-{5-[1-(Propane-1-sulfonyl) piperidin-4-yloxymethyl]-[1,2,4]oxadiazol-3-yl}pyridine;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid *tert*-butylamide;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidine-1-carboxylic acid *o*-tolylamide;
trans-4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)cyclohexanecarboxylic acid propyl ester;
trans-4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)cyclohexanecarboxylic acid butyl ester;
trans-4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-yl)cyclohexanecarboxylic acid isobutyl ester;
trans-4-[5-(4-Propoxymethylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine;
trans-4-[5-(4-Butoxymethylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine;
cis-4-[5-(3-Butoxymethylcyclopentyl)-[1,2,4]oxadiazol-3-yl]pyridine;
cis-4-[5-(3-Propoxymethylcyclopentyl)-[1,2,4]oxadiazol-3-yl]pyridine;
cis-4-[5-(3-Butoxymethylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)-3,4,5,6-tetrahydro-2H-[1,3']bipyridinyl;
2-[4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]pyrazine;
2-[4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]pyrimidine;
(4-Pentylcyclohexyl)-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amine;
(4-Pentylcyclohexyl-methyl)-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amine;
4-[(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;
4-[(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]methyl}-piperidine-1-carboxylic acid *tert*-butyl ester;
4-[[5-(2-Cyanopyridin-4-yl)-[1,2,4]oxadiazol-3-ylmethyl]amino]-piperidine-1-carboxylic acid *tert*-butyl ester
Methyl-(4-pentylcyclohexyl)-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amine;
Methyl-(4-pentylcyclohexylmethyl)-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amine;
4-[Methyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;
4-[Ethyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;
4-[Propyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;

4-[Cyclopropylmethyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;
4-[Butyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid *tert*-butyl ester;
4-{[Methyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]methyl}-piperidine-1-carboxylic acid *tert*-butyl ester;
4-{[Ethyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]methyl}-piperidine-1-carboxylic acid *tert*-butyl ester;
4-{[5-(2-Cyanopyridin-4-yl)-[1,2,4]oxadiazol-3-ylmethyl]ethylamino}-piperidine-1-carboxylic acid *tert*-butyl ester;
4-[Methyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid cyclopentyl ester;
4-[Methyl-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)amino]piperidine-1-carboxylic acid cyclopentyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxymethyl)piperidine-1-carboxylic acid *tert*-butyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethyl)piperazine-1-carboxylic acid *tert*-butyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethylsulfanyl)piperidine-1-carboxylic acid *tert*-butyl ester;
4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethanesulfonyl)piperidine-1-carboxylic acid *tert*-butyl ester
3-Pyridin-4-yl-[1,2,4]oxadiazole-5-carboxylic acid (4-pentylcyclohexyl)amide;
[4-(3-Pyridin-4-yl-[1,2,4]oxadiazol-5-ylmethoxy)piperidin-1-yl]phosphonic acid diphenyl ester;
4-{5-[2-(2H-Tetrazol-5-yl)pyridin-4-yl]-[1,2,4]oxadiazol-3-ylmethoxy}-piperidine-1-carboxylic acid *tert*-butyl ester;
4-[5-(2-Cyanopyridin-4-yl)-[1,2,4]oxadiazol-3-ylmethoxy]piperidine-1-carboxylic acid isopropyl ester;
and
4-[5-(2-Cyanopyridin-4-yl)-[1,2,4]oxadiazol-3-ylmethoxy]piperidine-1-carboxylic acid phenyl ester;
 or a pharmaceutically acceptable salt thereof.

16. **(Currently Amended)** A compound according to claim 1, or a pharmaceutically acceptable salt thereof, wherein B is $-\text{CH}=\text{CH}-$ or $(\text{CH}_2)_n$, where one of the CH_2 groups may be replaced by O, NR^5 , $\text{S}(\text{O})_m$ or $\text{C}(\text{O})$;

n is 2 or 3;

m is 0, 1 or 2;

R^2 is [[a]] 4- to 7-membered heterocyclyl containing one nitrogen atom which is substituted by $\text{C}(\text{O})\text{OR}^4$ or a 6-membered nitrogen containing heteroaryl group;

R^4 is C_{2-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, $\text{CO}_2\text{C}_{1-4}$ alkyl, $\text{N}(\text{R}^6)_2$ and NO_2 ;

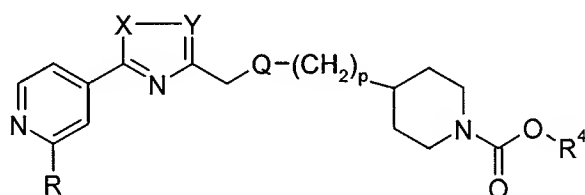
R^5 is hydrogen or C_{1-4} alkyl;

R^6 are independently hydrogen, or C_{1-4} alkyl, C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^9 , CN, SO_2CH_3 , $N(R^{10})_2$ and NO_2 ; or a group $N(R^{10})_2$ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR^{10} ;

R^9 is hydrogen, C_{1-2} alkyl or C_{1-2} fluoroalkyl; and

R^{10} is hydrogen or C_{1-4} alkyl.

17. **(Currently Amended)** A compound according to claim 1 having the formula (Ie), or a pharmaceutically acceptable salt thereof:



(Ie)

wherein one of X and Y is N, and the other is O;

Q is O, NR^5 or CH_2 ;

R is hydrogen, halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{3-7} cycloalkyl, aryl, OR^6 , CN, NO_2 , $S(O)_mR^6$, $CON(R^6)_2$, $N(R^6)_2$, $NR^{10}COR^6$, $NR^{10}SO_2R^6$, $SO_2N(R^6)_2$, a 4- to 7-membered heterocyclyl group or a 5- or 6-membered heteroaryl group;

R^4 is C_{2-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl, C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, CO_2C_{1-4} alkyl, $N(R^6)_2$ and NO_2 ;

R^5 is C_{1-4} alkyl;

R^6 are independently hydrogen, or C_{1-4} alkyl, C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^9 , CN, SO_2CH_3 , $N(R^{10})_2$ and NO_2 ; or a group $N(R^{10})_2$ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR^{10} ;

R^9 is hydrogen, C_{1-2} alkyl or C_{1-2} fluoroalkyl;

R^{10} is hydrogen or C_{1-4} alkyl; and

p is 0 or 1.

18. **(Previously Presented)** A pharmaceutical composition comprising a compound according to claim 1, including the compound of proviso c), or a pharmaceutically acceptable salt thereof; and a pharmaceutically acceptable carrier.

19. **(Previously Presented; Withdrawn)** A method for the treatment of a disease or condition in which GPR116 plays a role comprising a step of administering to a subject in need thereof an effective amount of a compound according to claim 1, including the compounds of provisos a) to c), or a pharmaceutically acceptable salt thereof.

20. **(Previously Presented; Withdrawn)** A method for the regulation of satiety comprising a step of administering to a subject in need thereof an effective amount of a compound according to claim 1, including the compounds of provisos a) to c), or a pharmaceutically acceptable salt thereof.

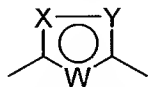
21. **(Previously Presented; Withdrawn)** A method for the treatment of obesity comprising a step of administering to a subject in need thereof an effective amount of a compound according to claim 1, including the compounds of provisos a) to c), or a pharmaceutically acceptable salt thereof.

22. **(Previously Presented; Withdrawn)** A method for the treatment of diabetes comprising a step of administering to a subject in need thereof an effective amount of a compound according to claim 1, including the compounds of provisos a) to c), or a pharmaceutically acceptable salt thereof.

23. **(Currently Amended; Withdrawn)** A method for the treatment of a disease or condition in which GPR116 plays a role comprising a step of administering to a subject in need thereof an effective amount of a compound of the formula:



or a pharmaceutically acceptable salt thereof;
wherein V represents a 5-membered heteroaryl ring of the formula:



wherein W is N and one of X and Y is N and the other is O;

B is -CH=CH- or (CH₂)_n, where one of the CH₂ groups may be replaced by O, NR⁵, S(O)_m, C(O) or C(O)NR¹²;

n is 0, 1, 2 or 3;

m is 0, 1 or 2;

R^1 is 3- or 4-pyridyl, 4- or 5-pyrimidinyl or 2-pyrazinyl, any of which may be optionally substituted by one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{3-7} cycloalkyl, aryl, OR^6 , CN, NO_2 , $S(O)_mR^6$, $CON(R^6)_2$, $N(R^6)_2$, $NR^{10}COR^6$, $NR^{10}SO_2R^6$, $SO_2N(R^6)_2$, a 4- to 7-membered heterocyclyl group or a 5- or 6-membered heteroaryl group;

R^2 is 4- to 7-membered cycloalkyl substituted by R^3 , $C(O)OR^3$, $C(O)R^3$ or $S(O)_2R^3$, or R^2 is 4- to 7-membered heterocyclyl, wherein the heterocycle contains one nitrogen atom which is substituted by containing one or two nitrogen atoms which is unsubstituted or substituted by $C(O)OR^4$, $C(O)R^3$, $S(O)_2R^3$, $C(O)NHR^4$, $P(O)(OR^{11})_2$ or a 5- or 6-membered nitrogen containing heteroaryl group;

R^3 is C_{3-8} alkyl, C_{3-8} alkenyl or C_{3-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be optionally substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, CO_2C_{1-4} alkyl, $N(R^6)_2$ and NO_2 ;

R^4 is C_{2-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, CO_2C_{1-4} alkyl, $N(R^6)_2$ and NO_2 ;

R^5 is hydrogen, $C(O)R^7$, $S(O)_2R^8$, C_{3-7} cycloalkyl or C_{1-4} alkyl optionally substituted by OR^6 , C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-2} alkyl, C_{1-2} fluoroalkyl, OR^6 , CN, $N(R^6)_2$ and NO_2 ;

R^6 are independently hydrogen C_{1-4} alkyl, C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^9 , CN, SO_2CH_3 , $N(R^{10})_2$ and NO_2 ; or a group $N(R^{10})_2$ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR^{10} ;

R^7 is hydrogen, C_{1-4} alkyl, OR^6 , $N(R^6)_2$, aryl or heteroaryl;

R^8 is C_{1-4} alkyl, C_{1-4} fluoroalkyl, aryl or heteroaryl;

R^9 is hydrogen, C_{1-2} alkyl or C_{1-2} fluoroalkyl;

R^{10} is hydrogen or C_{1-4} alkyl;

R^{11} is phenyl; and

R^{12} is hydrogen, C_{1-4} alkyl or C_{3-7} cycloalkyl.